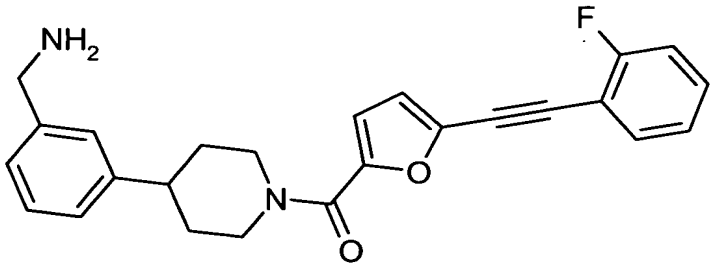
WHAT IS CLAIMED IS:

1. A compound of Formula (I):



(I)

or a prodrug, pharmaceutically acceptable salt, or solvate thereof.

2. The compound of Claim 1 as a pharmaceutically acceptable salt thereof.

3. The compound of Claim 2 wherein the pharmaceutically acceptable salt is a trifluoroacetate or methanesulfonate salt.

4. A method for treating a patient suffering from, or subject to, a physiological condition in need of amelioration comprising administering to the patient a therapeutically effective amount of the compound of Claim 1.

5. The method of Claim 4, wherein the physiological condition is selected from the group consisting of inflammatory disease, a disease of joint cartilage destruction, ocular conjunctivitis, vernal conjunctivitis, inflammatory bowel disease, asthma, allergic rhinitis, interstitial lung disease, fibrosis, sceleroderma, pulmonary fibrosis, liver cirrhosis, myocardial fibrosis, neurofibroma, hypertrophic scar, dermatological condition, condition related to atherosclerotic plaque rupture, periodontal disease, diabetic retinopathy, tumor growth, anaphylaxis, multiple sclerosis, peptic ulcer, and syncytial viral infection.

6. The method of Claim 5, wherein the physiological condition is inflammatory disease.

7. The method of Claim 6 wherein the inflammatory disease is joint inflammation, arthritis, rheumatoid arthritis, rheumatoid spondylitis, gouty arthritis, traumatic arthritis, rubella arthritis, psoriatic arthritis, or osteoarthritis.

8. The method of Claim 5, wherein the physiological condition is a dermatological condition.

9. The method of Claim 8, wherein the dermatological condition is atopic dermatitis or psoriasis.

10. The method of Claim 5, wherein the physiological condition is related to atherosclerotic plaque rupture.

11. The method of Claim 10, wherein the physiological condition related to atherosclerotic plaque rupture is myocardial infarction, stroke, or angina.

12. A method for treating a patient suffering from asthma, comprising administering to the

patient a combination of a thereapeutically effective amount of a compound of Claim 1, and a second compound selected from the group consisting of a beta andrenergic agonist, anticholinergic, anti-inflammatory corticosteroid, and anti-inflammatory agent.

13. A pharmaceutical composition comprising a therapeutically effective amount of a compound of Claim 1 and a pharmaceutically acceptable carrier thereof.

14. A pharmaceutical composition comprising a compound of Claim 1 and a therapeutically effective amount of a second compound selected from the group consisting of a beta andrenergic agonist, anticholinergic, anti-inflammatory corticosteroid, and anti-inflammatory agent; and a pharmaceutically acceptable carrier.

15. The pharmaceutical composition of Claim 14, wherein the second compound is a beta andrenergic agonist.

16. The pharmaceutical composition of Claim 15, wherein the beta andrenergic agonist is selected from albuterol, terbutaline, formoterol, fenoterol, or prenaline.

17. The pharmaceutical composition of Claim 14, wherein the second compound is an anticholinergic.

18. The pharmaceutical composition of Claim 17, wherein the anticholinergic is ipratropium bromide.

19. The pharmaceutical composition of Claim 14, wherein the second compound is an anti-inflammatory corticosteroid.

20. The pharmaceutical composition of Claim 19, wherein the anti-inflammatory corticosteroid is selected from beclomethasone dipropionate, triamcinolone acetonide, flunisolide or dexamethasone.

21. The pharmaceutical composition of Claim 14, wherein the second compound is an anti-inflammatory agent.

22. The pharmaceutical composition of Claim 21, wherein the anti-inflammatory agent is soldium cromoglycate or nedocromil sodium.

23. The pharmaceutical composition of Claim 14, wherein the second compound is a pharmaceutically acceptable carrier thereof.